Please amend the following claims:

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26. (currently amended) A method for treating a patient suffering from [preventing and treating] the side-effects of a ketogenic diet, said method comprising the steps of:

administering a composition of a plurality of agents that produces a synergistic effect of reducing the concentration of a plurality of internal body chemicals, wherein said composition is comprised of,

a hypocholesterolemic agent, wherein said hypocholesterolemic agent is selected from the group consisting of benfluorex and ursodesoxycolic acid;

a hypotriglyceride agent, wherein said hypotriglyceride agent is benfluorex;

a lipasic and proteasic agent, wherein said lipasic and proteasic agent is pancreatine IX F.U.;

a hypoglycemic agent, wherein said hypoglycemic agent is metformine; and

a hydrocoleretic agent, wherein said hydrocoleretic agent is selected from the group consisting of Na dehydrocloatye and ursodesoxycolic acid.

27. (currently amended) The method as claimed in claim [1] 26, wherein in said administration of said composition, said composition further comprises at least one of:

a hypouricemic agent, wherein said hypouricemic agent is centella asiatica purified [trierpenes]triterpenes;

a radical scavenger agent, wherin said radical scavenger agent is selenium;

a sympatholytic agent, wherein said sympatholytic agent is [yohinbine] yohimbine;

a sympathicomimetic agent, wherein said sympathicomimetic agent is from the group consisting of [phendimetrazinum] phendimetrazine bitartrate and phendimetrazinum pamoate; and

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at least one vitamin, wherein said at least on vitamin being selected from the group consisting of vitamin A, vitamin B1, vitamin B6, vitamin E and Vitamin C.

28. (previously added)The method as claimed in claim 27, wherein in said administration of said composition, said composition further comprises at least one diet adjuvant selected from the group consisting of sedative-ansiolytic agents, anoretic agents and lipolytic agents.

29. (currently amended) The method as claimed in claim 26, wherein in said administration of said composition, said benfluorex is present in [global] amount from 7% to 23% in weight of the total amount of the composition;

said pancreatine IX F.U. is present in an amount from 27% to 43% in weight of the total amount of the composition;

said metformine is present in an amount from 36% to 41% in weight of the total amount of the composition; and

said Na dehydrocloate is present in an amount from 9% to 14% of the total amount of the composition.

30. (previously added) The method as claimed in claim 26, wherein in said administration of said composition,

said benfluorex is present in an amount from 7% to 23% in weight of the total amount of the composition;

said pancreatine IX F.U. is present in an amount from 27% to 43% in weight of the total amount of the composition;

said metformine is present in an amount from 36% to 41% in weight of the total amount of the composition; and

said Ursodesoxycolic acid is present in an amount from 14% to 17% in weight of the total amount of the composition.

31. (currently amended) The method as claimed in claim 27, wherein in said administration of said composition, each of said elements of said composition are in a ratio of weight with respect to the total weight of said composition [at least one of a set of ratios, said ratios including the set of] wherein,

said centella asiatica purified [trierpenes] triterpenes is in a ratio from 0.04:1 to 0.5:1 in weight with respect to [the] said total weight of composition;

said selenium is in a ratio from 0.09:1 to 0.3:1 in weight with respect to [the] said total weight of composition;

said yohimbine is in a ratio from 0.0009:1 to 0.0007:1 in weight with respect to [the] said total weight of composition;

said phendimetrazine bitartarate or phendimetrazine pamoate is in

a ratio from 0.004:1 to 0.13:1 in weight with respect to [the] said total weight of composition;

said vitamin A is in a ratio from 0.5:1 to 1.8:1 in weight with respect to [the] said total weight of composition;

said vitamin B1, is in a ratio from 0.002:1 to 0.2:1 in weight with respect to [the] said total weight of composition;

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said vitamin B6, is in a ratio from 0.05:1 to 0.2:1 in weight with respect to [the] said total weight of composition;

said vitamin E, is in a ratio from 0.09:1 to 1:1 in weight with respect to [the] said total weight of composition; and

said vitamin C, is in a ratio from 0.09:1 to 0.3:1 in weight with respect to the total weight of composition.

32. (currently amended) The method as claim in claim 28, wherein in said administration of said composition, each of said elements of said composition are in a ratio of weight with respect to the total weight of the composition [at least one of a set of ratios, said ratios including the set of] wherein ;

said sedative-ansiolityc agent is the benzodiazepine dipotassium chlorazepate in a ratio from 0.0005:1 to 0.03:1 in weight with respect to the total weight of composition;

said anoretic agent is selected from the group consisting of diethylpropione chlorhydrate, fenfluramine chlorhydrate, D-fenfluramine chlorhydrate, said anorectic agent being present in a ratio from 0.002:1 to 1.3:1 in weight with respect

to [the] said total weight of composition; and

said lipolityc agent is selected from the group consisting of [the analogue of tiroxine,] triiodiotiroacetic acid and any tiroxine analogs which is present in a ratio from 0.0002:1 to 0.003:1 in weight with respect to [the] said total weigh of composition.

33. (previously added) The method as claimed in claim 26, wherein said administration of said composition is performed orally in doses from 7g to 23g per day to a patient of the weight of approximately 70kg.

34. (previously added)The method as claimed in claim 27, wherein said administration of said composition is performed orally in doses from 7g to 23g per day to a patient of the weight of approximately 70kg.

35. (previously added)The method as claimed in claim 28, wherein said administration of said composition is performed orally in doses from 7g to 23g per day to a patient of the weight of approximately 70kg.

36. (previously added)The method as claimed in claim 29, wherein said administration of said composition is performed orally in doses from 7g to 23g per day to a patient of the weight of approximately 70kg.

37. (previously added)The method as claimed in claim 30, wherein said administration of said composition is performed orally in doses from 7g to 23g per day to

a patient of the weight of approximately 70kg.

38. (previously added)The method as claimed in claim 31, wherein said administration of said composition is performed orally in doses from 7g to 23g per day to a patient of the weight of approximately 70kg.

39. (previously added)The method as claimed in claim 32, wherein said administration of said composition is performed orally in doses from 7g to 23g per day to a patient of the weight of approximately 70kg.

40. (currently added) he method as claimed in claim 26, wherein in said administration of said composition, further comprises the administering of said composition to reduce fibrinogen levels.